

AMENDMENTS TO THE CLAIMS

Please amend the claims without prejudice, without admission, without surrender of subject matter, and without any intention of creating any estoppel as to equivalents, as follows:

1. (Currently amended) A stable formulation suitable for administration to animals comprising a combination of two active ingredients and a pyrrolidone solvent, wherein the said combination of two active ingredients consists of levamisole and an avermectin or levamisole and a milbemycin at least one active selected from the group consisting of avermectins and milbemycins and levamisole and both of said actives being dissolved in a pyrrolidone solvent.
2. (Previously presented) The stable formulation suitable for administration to animals as claimed in claim 1, additionally including a co-solvent selected from the group consisting of glycol ethers.
3. (Original) The stable formulation suitable for administration to animals as claimed in claim 1, wherein the pyrrolidone solvent is 2-pyrrolidone or N-methyl pyrrolidone.
4. (Original) The stable formulation suitable for administration to animals as claimed in claim 1, wherein the avermectin or milbemycin is present in the range of between 0.01-5% w/v.
5. (Previously presented) The stable formulation suitable for administration to animals as claimed in claim 4, wherein the avermectin or milbemycin is selected from the group consisting of abamectin, doramectin, eprinomectin, ivermectin and moxidectin.
6. (Original) The stable formulation suitable for administration to animals as claimed in claim 1, wherein the levamisole is present in the range of between 1-30% w/v.
7. (Currently amended) A stable formulation suitable administration to animals as claimed in claim 1, wherein the formulation additionally includes at least one further medicament selected from the group consisting of anthelmintics, dietary supplements, vitamins, mineral, preservatives, stabilisers, flavorants, co-solvents and other inactive excipients beneficial agents.
8. (Original) The stable formulation suitable for administration to animals as claimed in claim 1, wherein the formulation is suitable for topical administration.

9. (Original) The stable formulation suitable for administration to animals as claimed in claim 1, wherein the formulation is suitable for parenteral administration.

10. (Original) The stable formulation suitable for administration to animals as claimed in claim 1, wherein the formulation is suitable for oral administration.

11. (Currently amended) A method of treating ~~or preventing infection of cattle with cattle infected with~~ Cooperia or Ostertagia by administering a formulation as claimed in claim 1.